Appl. No. 09/932,538 Amdt. dated January 24, 2005 Reply to Office Action of August 24, 2004 Page 7 of 12

## **Amendments to the Drawings**

Please re-include Figure 8 as the last page of the drawings, a copy of which is attached.

Attachment:

Replacement Sheet

**REMARKS/ARGUMENTS** 

I. Introduction

Upon entry of the present amendment, claims 5-7 remain under examination in this

application. Figure 8 and its accompanying description have been re-included in the

application. No new matter has been added. Also with the response, Applicant submits a

declaration from Dr. Relyveld showing a distinction between the material, structural, and

functional characteristics of the claimed composition and the composition of the cited prior

art – U.S. Patent No. 4,016,252 to Relyveld.

Because the present amendments (1) do not raise new issues requiring further

consideration or search, (2) do not introduce new matter, (3) materially reduce the issues for

appeal, and (4) place this application into better condition for allowance, entry is appropriate

under 37 C.F.R. § 1.116, and is respectfully requested. Applicant respectfully requests

reconsideration and allowance of the pending claims.

II. Amendments to the Specification and the Drawings

The Examiner originally objected to Figure 8, stating that "the specification fails to

describe Figure 8." See Office action dated April 7, 2004. Without acquiescing to

Examiner's reasoning, Applicant cancelled Figure 8. See Response filed June 8, 2004. (One

reason for this cancellation was because the ocular embodiment of this invention is being

claimed in a related pending application.)

ATLLIB01 1910159.1

Appl. No. 09/932,538

Amdt. dated January 24, 2005

Reply to Office Action of August 24, 2004

Page 9 of 12

However, now the Examiner's position is that deletion of Figure 8 constitutes new

matter because the figure provided support for enablement of the claims. This is a confusing

rejection (and one in which the Applicant does not agree), but nonetheless, Figure 8 has been

re-included in the application. Applicant is also re-including the description of Figure 8.

Applicant submits that the amendments to the specification and Figure 8 remove the grounds

for Examiner's rejection, and respectfully request that it be withdrawn.

III. 35 U.S.C. §112 Rejections

The Examiner has rejected claims 5-7 under 35 U.S.C. §112, second paragraph, as

being indefinite. The Examiner's position is that claim 5 is rendered vague and indefinite by

the use of the phrase: "induce immunity in a patient."

Applicant continues to submit that the phrase, by its plain and ordinary meaning,

means to induce an immunological response in a patient. Applicant refers the Examiner to

the following portions of the specification, which discuss possible immunological responses

that are sought to be induced:

• cell mediated immunity and antibody responses (page 4, lines 18-24);

• strong Th1 T-cell associated enhancement of microbial immunity, especially IgA

production and anti-viral cell-mediated immunity (CMI) (page 5, lines 15-19);

• mucosal IgA immunity (page 5, lines 20-25);

• humoral and cell-mediated immunity, with the mucosal immunity manifested as a

humoral response (page 8, line 25 - 26);

• stimulation of B cells which differentiate into antibody-producing plasma cells;

antibodies are capable of recognizing extracellular pathogens, while the cell-mediated

ATLLIB01 1910159.1

Appl. No. 09/932,538

Amdt. dated January 24, 2005

Reply to Office Action of August 24, 2004

Page 10 of 12

component involves T lymphocytes capable of recognizing intracellular pathogens (page 8, line 24 – page 9, line 12);

- systemic and mucosal immunity, with relative absence of side effects and lack of IgE antibody production (page13, lines 5-10);
- increased survival rates and less severe clinical infection compared to controls, following live-viral challenge (page 13, lines 14-19);
- antigen-specific immunity (page 14, lines 22-25);
- induce higher IgG titers and IgA levels (page 35, lines 16-21).

Based on all of this disclosure in the specification, coupled with the understanding that one of ordinary skill in the art would have, Applicant respectfully submits that the phrase "induce immunity in a patient" is not indefinite or vague. One of ordinary skill in the art would understand the immunity that is sought to be induced by the present invention, particularly when the claim is read in light of the specification. Accordingly, Applicant respectfully requests that the rejection be withdrawn.

## IV. 35 U.S.C. §102 Rejection

The Examiner has also maintained the rejection of claims 5-7 under 35 U.S.C. §102(b) as being anticipated by Relyveld (U.S. Patent 4,016,252). Applicant traverses this rejection and respectfully requests reconsideration and withdrawal thereof.

Applicant submits with this response a declaration by Dr. Relyveld dated November 18, 2004, in which Dr. Relyveld states that "...I have compared the calcium phosphate particles disclosed in the Application to those disclosed in U.S. Patent 4,016,252 ("the '252 patent"), issued to me." *See* attached declaration of Dr. Relyveld, ¶4. "The calcium

Appl. No. 09/932,538

Amdt. dated January 24, 2005

Reply to Office Action of August 24, 2004

Page 11 of 12

phosphate particles disclosed in the Application are different from the calcium phosphate

particles disclosed in the '252 Patent." ¶6. "Unlike the present invention, the '252 Patent

does not disclose calcium phosphate particles that are 'substantially smooth' or 'substantially

spherical.' Despite efforts, I was unable to obtain the 'substantially smooth' or 'substantially

spherical' calcium phosphate particles of the present invention. The inventors of the

Application were able to obtain these unexpected results through their novel manufacturing

technique." ¶7. "The novel morphology of the calcium phosphate particles of the present

invention result in greater control over the degree of antigenic material saturation which can

be achieved in the particle. This control leads to greater efficiency in particle production and

efficacy in treatment using the particles." ¶8. "The Action is incorrect in stating that the

particles in the '252 Patent are 'of the same composition and the same size' as the calcium

phosphate particles of the present invention." ¶ 9.

Because Applicant has met its burden of "show[ing] a distinction between the

material, structural and functional characteristics of the claimed composition and the

composition of the prior art," Applicant respectfully requests withdrawal of this rejection

and prompt issuance of a patent containing the pending claims.

Attachment: Relyveld Declaration

ATLLIB01 1910159.1

Appl. No. 09/932,538 Amdt. dated January 24, 2005 Reply to Office Action of August 24, 2004 Page 12 of 13

## **CONCLUSION**

Applicant respectfully submits that claims 5-7 are in condition for immediate allowance, and requests early notification to that effect. If any issues remain to be resolved, the Examiner is respectfully requested to contact the undersigned at 404.815.6147 to arrange for a telephone interview.

Respectfully submitted,

Kristin M. Crall Reg. No. 46,895

KILPATRICK STOCKTON LLP 1100 Peachtree Street Suite 2800 Atlanta, Georgia, 30309-4530 404.815.6147